

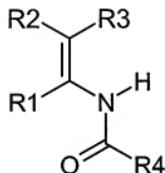
Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims:

1-11. (Cancelled)

12. (New) A process for the production of enamide derivatives represented by formula (I)



(I)

wherein;

R1 and R2 and R3 are independently selected from the group consisting of a hydrogen atom; an alkyl; a cycloalkyl; a cycloalkylalkyl; an alkylaryl; an aryl; a heterocycle; a cyano; an alkoxy; an aryloxy; a carboxyl; a carbamoyl; -CONRSR6 in which R5 and R6 are independently selected from an alkyl, an arylalkyl, an aryl; and R5 and R6 taken together may form a ring; and -COOR5 in which R5 is selected from an alkyl, an alkylaryl, a cycloalkyl, and aryl;

said alkyl, cycloalkyl, cycloalkylalkyl, alkylaryl and aryl being substituted or not substituted with a group selected from a functional group and R5;

R1 and R2 taken together may form a monocyclic ring ; a di-cyclic ring and a higher polycyclic ring, said ring being substituted or not substituted with a group selected from a functional group and R5;

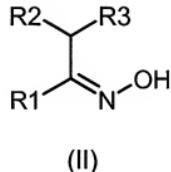
R4 is selected from the group consisting of a hydrogen atom, alkyl, aryl and alkylaryl; said alkyl, aryl, and alkylaryl being substituted or not substituted with halogen;

X is selected from an oxygen atom or a leaving group;

m is an integer selected from 1 and 2;

when m is 1 X is a leaving group; when m is 2 X is an oxygen atom;

said method comprising a hydrogenation/isomerization reaction in presence of a heterogeneous catalyst based on at least one metal selected from Pd, Ir, Pt, Rh, or Ni, of an oxime derivative of formula (II)



wherein R1, R2 and R3 are as defined above;

with an acyl derivative of formula (III):



wherein R4, m and X are as defined above.

13. (New) The process of claims 12, wherein the derivative of formula (III) is used in the amount selected from at least 2 times per mole based on the oxime, and an amount sufficient to act as a reacting agent and as a solvent.

14. (New) The process of claim 12, wherein the heterogeneous catalyst is a metal is selected from the group consisting of Ir and Rh.

15. (New) The process of claim 12, wherein the heterogeneous catalyst is in a form selected from a metal oxide and from a metallic form, optionally supported on a suitable carrier; and is used in an amount ranging between 0.001 and 30% mole, based on the oxime derivative.

16. (New) The process of claim 12, which is carried out in a suitable solvent.

17. (New) The process of claim 12, which is carried out under a hydrogen pressure ranging between 0.5 and 20 bars.

18. (New) The process of claim 12, which is carried out under a temperature ranging between -20 and 150 °C.

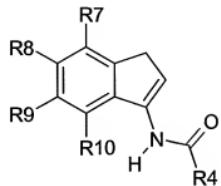
19. (New) The process of claim 12, further comprising a work up step of an organic solution of the compound of formula (I) which is a washing step with water containing organic or mineral salt(s) without halogen atom.

20. (New) The process of claim 19, wherein the organic or mineral salt(s) is/ are selected from the group consisting of phosphate, sulfate, acetate, citrate, formate, borate, carbonate, or ammonium.

21. (New) The process of claim 12, wherein said eneamide is selected from the group consisting of:

- N-(6-Methoxy-3H-inden-1-yl)-acetamide;
- N(3,4-dihydro-1-naphthalenyl)acetamide;
- N(3,4-dihydro-naphthalen-2-yl)acetamide;
- N-(2-Phenyl-cyclohex-1-enyl)-acetamide; and
- N-(7-Methoxy-3,4-dihydro-naphthalen-2-yl)-acetamide.

22. (New) A method of manufacture of an amine or an amide compound aimed in the preparation of a pharmaceutical substance comprising performing a hydrogenation reaction of a eneamide compound of formula (IIE)



wherein R4 is selected from the group consisting of hydrogen, alkyl, aryl and alkylaryl; said alkyl, aryl, and alkylaryl being substituted or not substituted with halogen;

R7, R8, R9 and R10 are independently selected from the group consisting of hydrogen, functional group, alkyl and aryl, while not simultaneously being hydrogen;

to obtain a hydrogenated compound; and

said hydrogenated compound of formula (IIE) is further used as an intermediate in the synthesis of said pharmaceutical substance.

23. (New) The method of claim 22, wherein said hydrogenation reaction performs an asymmetric hydrogenation of said compound of formula (IIE).

24. (New) The method of claim 22, wherein the eneamide compound is selected from the group consisting of:

- N-(6-Methoxy-3H-inden-1-yl)-acetamide;
- N(3,4-dihydro-1-naphthalenyl)acetamide;
- N(3,4-dihydro-naphthalen-2-yl)acetamide;
- N-(2-Phenyl-cyclohex-1-enyl)-acetamide; and
- N-(7-Methoxy-3,4-dihydro-naphthalen-2-yl)-acetamide.